

Amendments to the Claim:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1-7. (Cancelled)

8. (Currently Amended) A purified compound capable of binding to the neural cell adhesion molecule (NCAM) homophylic binding site composed of the Ig1, Ig2 and Ig3 modules, wherein said compound is capable of

- i) binding to the Ig1 module of NCAM at said NCAM homophylic binding site, and thereby mimicking and/or modulating the interaction between the Ig1 and Ig3 modules of NCAM, wherein said modules are from two individual NCAM molecules of opposing contacting cells, and/or
- ii) binding to the Ig3 module of NCAM at said NCAM homophylic binding site, and thereby mimicking and/or modulating the interaction between the Ig3 and Ig1 modules of NCAM, wherein said modules are from two individual NCAM molecules of opposing contacting cells, and/or
- iii) binding to the Ig2 module of NCAM at said NCAM homophylic binding site, and thereby mimicking the interaction between Ig2 and Ig3 modules of NCAM, wherein said modules are from two individual NCAM molecules of opposing contacting cells, and/or
- iv) binding to the Ig3 module of NCAM at said NCAM homophylic binding site, and thereby mimicking and/or modulating the binding between the Ig3 and Ig2 modules of NCAM, wherein said modules are from two individual NCAM molecules of opposing contacting cells, and/or
- v) binding to the Ig2 module of NCAM at said NCAM homophylic binding site, and thereby mimicking and/or modulating the interaction between the Ig2 and Ig2 modules of NCAM, wherein said modules are from two individual NCAM

molecules of opposing contacting cells,
said compound being

(I) (a) a peptide sequence identified as SEQ ID NO: 1, 2,
4, 6, 7, 8, 9, 10, 11, 12, 13, 14, 16, 17, 18, 40 or 41,

(b) a peptide which is a fragment of (a), consisting of
at least 5 amino acids,

(c) a peptide consisting of a peptide sequence according
to (a) and up to 10 additional amino acids,

(d) a peptide which differs from (a), (b) or (c) solely
by one or more amino acid substitutions, but (d1) comprises at
least a five amino acid fragment of a peptide of (a) or (d2)
comprises a sequence at least 50% 90% identical to a peptide
of (a), or

(e) a peptide sequence identified as SEQ ID NO:5; or

(II) an oligomer or polymer comprising a plurality of
peptides according to (I) above, which oligomer or polymer
either consists of a plurality of peptides according to (I)
above, or comprises a non-NCAM carrier moiety to which said
peptides are attached.

9. (Previously Presented) The compound according to claim
8, said compound consisting of the amino acid sequence
WFSPNNGEKLSQPNQ (SEQ ID NO: 1).

10. (Previously Presented) The compound according to
claim 8, said compound consisting of the amino acid sequence
YKCVVTAEDGTQSE (SEQ ID NO: 2).

11. (Cancelled).

12. (Previously Presented) The compound according to
claim 8, said compound consisting of the amino acid sequence
QIRGIKKTD (SEQ ID NO: 4).

13. (Previously Presented) The compound according to
claim 8, said compound consisting of the amino acid sequence
DVR (SEQ ID NO: 5).

14. (Previously Presented) The compound according to
claim 8, said compound consisting of the amino acid sequence

RGIKKTD (SEQ ID NO: 6).

15. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence DVRRGIKKTD (SEQ ID NO: 7).

16. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence KEGED (SEQ ID NO: 8).

17. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence IRGIKKTD (SEQ ID NO: 9).

18. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence KEGEDGIRGIKKTD (SEQ ID NO: 10).

19. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence DKNDE (SEQ ID NO: 11).

20. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence TVQARNSIVNAT (SEQ ID NO: 12).

21. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence SIHLKVFVAK (SEQ ID NO: 13).

22. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence LSNNYLQIR (SEQ ID NO: 14).

23. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence RFIVLSNNYLQI (SEQ ID NO: 15).

24. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence KKDVRFIVLSNNYLQI (SEQ ID NO: 16).

25. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence QEFKEGEDAVIV (SEQ ID NO: 17).

26. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence KEGEDAVIVCD (SEQ ID NO: 18).

27. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence AFSPNGEKLSPNQ (SEQ ID NO: 40).

28. (Previously Presented) The compound according to claim 8, said compound consisting of the amino acid sequence AKSVVTAEDGTQSE (SEQ ID NO: 41).

29. (Cancelled)

30. (Withdrawn) The method of claim 42, wherein the medicament is for treating normal, degenerated or damaged NCAM presenting cells.

31. (Withdrawn) The method of claim 42, wherein the medicament is for treatment comprising the stimulation of differentiation and/or survival of NCAM presenting cells.

32. (Withdrawn) The method of claim 42, wherein the medicament is for treating the diseases and conditions of the central and peripheral nervous system, or of the muscles or of various organs.

33. (Withdrawn) The method of claim 42, wherein the medicament is for treating the diseases or conditions of the central and peripheral nervous system, such as postoperative nerve damage, traumatic nerve damage, impaired myelination of nerve fibers, postischaemic damage, e.g. resulting from a stroke, Parkinson's disease, Alzheimer's disease, Huntington's disease, dementias such as multiinfarct dementia, sclerosis, nerve degeneration associated with diabetes mellitus, disorders affecting the circadian clock or neuro-muscular transmission, and schizophrenia, mood disorders, such as manic depression; for treatment of diseases or conditions of the muscles including conditions with impaired function of neuro-muscular connections, such as after organ transplantation, or such as genetic or traumatic atrophic

muscle disorders; or for treatment of diseases or conditions of various organs, such as degenerative conditions of the gonads, of the pancreas such as diabetes mellitus type I and II, of the kidney such as nephrosis and of the heart, liver and bowel.

34. (Withdrawn) The method of claim 42, wherein the medicament is for treating the postoperative nerve damage, traumatic nerve damage, impaired myelination of nerve fibers, postischaemic, e.g. resulting from a stroke, Parkinson's disease, Alzheimer's disease, dementias such as multiinfarct dementia, sclerosis, nerve degeneration associated with diabetes mellitus, disorders affecting the circadian clock or neuro-muscular transmission, and schizophrenia, mood disorders, such as manic depression.

35. (Withdrawn) The method of claim 42, wherein the medicament is for promoting the wound-healing.

36. (Withdrawn) The method of claim 42, wherein the medicament is for treating the cancer.

37. (Withdrawn) The method of claim 42, wherein the medicament is for preventing the cell death of heart muscle cells, such as after acute myocardial infarction, or after angiogenesis.

38. (Withdrawn) The method of claim 42, wherein the medicament is for promoting the revascularisation.

39. (Withdrawn) The method of claim 42, wherein the medicament is for stimulating the ability to learn and/or of the short and/or long-term memory.

40. (Cancelled)

41. (Previously Presented) A pharmaceutical composition comprising one or more compounds as defined in claim 8.

42. (Withdrawn) A method of treating a disease wherein modulating differentiation, adhesion and/or survival of NCAM presenting cells is essential for the treatment, which comprises administering to a subject in need thereof a

therapeutically effective amount of a medicament comprising a compound according to claim 8.

43. (Cancelled)

44. (Currently Amended) The compound according to claim 8, wherein said peptide sequence is selected by a method comprising the steps of

- a) providing a crystalline polypeptide according to claim 1 comprising the Ig1-2-3 module of NCAM, said polypeptide comprising amino acid residues 1 to 289 of SEQ ID NO: 44, wherein said crystal comprises atoms arranged in a spatial relationship represented by the structure co-ordinates of Table 2 (Figure 2) or by coordinates having a root mean square deviation therefrom of not more than 2.5 Å,
- b) generating a structural model of the Ig1-2-3 module of NCAM of (a) by using the computer modelling techniques;
- c) in-silico evaluating compounds for the capability of
 - i) binding to the Ig1 module of NCAM at the NCAM homophytic binding site composed of the Ig1, Ig2 and Ig3 modules, and thereby mimicking and/or modulating the interaction between the Ig1 and Ig3 modules of NCAM, wherein said modules are from two individual NCAM molecules, and/or
 - ii) binding to the Ig3 module of NCAM at the NCAM homophytic binding site composed of the Ig1, Ig2 and Ig3 modules, and thereby mimicking and/or modulating the interaction between the Ig3 and Ig1 modules of NCAM, wherein said modules are from two individual NCAM molecules, and/or
 - iii) binding to the Ig2 module of NCAM at the NCAM homophytic binding site composed of the Ig1, Ig2 and

Ig3 modules, and thereby mimicking the interaction between Ig2 and Ig3 modules of NCAM, wherein said modules are from two individual NCAM molecules, and/or

- iv) binding to the Ig3 module of NCAM at the NCAM homophylic binding site composed of the Ig1, Ig2 and Ig3 modules, and thereby mimicking and/or modulating the binding between the Ig3 and Ig2 modules of NCAM, wherein said modules are from two individual NCAM molecules, and/or
- v) binding to the Ig2 module of NCAM at the NCAM homophylic binding site composed of the Ig1, Ig2 and Ig3 modules, and thereby mimicking and/or modulating the interaction between the Ig2 and Ig2 modules of NCAM, wherein said modules are from two individual NCAM molecules,

by using the structural model of the Ig1-2-3 module of NCAM of (b);

- d) selecting a candidate compound capable of at least one interaction of (c), and
- e) testing the candidate compound of (d) in an in vitro assay for the capability of modulating differentiation, adhesion and/or survival of NCAM presenting cells, said assays comprising at least one NCAM presenting cell, and /or

testing the candidate compound of (d) in an assay comprising evaluating the capability of the compound of at least one interaction of (b) by contacting the compound with at least one individual fragment of an NCAM molecule, said fragment comprising a sequence of consecutive amino acid residues corresponding to the sequence of the Ig1-2-3 module of NCAM comprising residues 1 to 289 of the sequence set forth in SEQ ID NO: 44.

45. (Previously Presented) The compound according to

claim 8 wherein the carrier of (II) is a lysine dendrimer backbone.

46. (Previously Presented) The compound according to claim 8 wherein the carrier of (II) is a protein.

47. (Previously Presented) The compound according to claim 46 wherein the carrier of (II) is bovine serum albumin.

48. (Previously Presented) The compound according to claim 8 wherein the carrier of (II) is a lipophilic dendrimer, a micelle-like carrier formed by lipophilic derivatives, a starburst carbon chain polymer conjugate, or a ligand presenting assembly based on a derivative of diethylaminomethane.

49. (Previously Presented) The compound according to claim 8 wherein (I) applies.

50. (Previously Presented) The compound of claim 49 wherein the peptide of (I) (d) comprises a sequence at least 85% identical to a peptide of (I) (a).

51. (Previously Presented) The compound of claim 49 wherein the peptide of (I) (d) comprises a sequence differing from a peptide of (I) (a) solely by a single substitution.

52. (Previously Presented) The compound of claim 49 wherein the peptide of (I) (d) comprises a sequence which differs from a peptide of (I) (a) solely by one or more conservative substitutions, such being defined as

(1) replacement of an amino acid selected from the group consisting of Gly, Ala, Val, Leu and Ile with another amino acid of the same group,

(2) replacement of an amino acid selected from group consisting of Asp, Glu, Asn and Gln with another amino acid of the same group,

(3) replacement of an amino acid selected from the group consisting of Phe, Tyr, Trp, His, and Pro with another amino acid of the same group,

(4) replacement of an amino acid selected from the group

consisting of Arg, Lys and His with another amino acid of the same group,

(5) replacement of Cys with an amino acid selected from the group consisting of Asp, Glu, Lys, Arg, His, Asn, Gln, Ser, Thr and Tyr,

(6) replacement of an amino acid selected from the group consisting of Pro, Ala, Gly, Ser and Tyr with another amino acid of the same group, and

(7) replacement of an amino acid selected from the group consisting of Leu, Ile, Val and Met with another amino acid of the same group.

53. (Previously Presented) The composition of claim 52 wherein the peptide of (I) (c) comprises a sequence differing from a peptide of (I) (a) solely by a single conservative substitution.

54. (Previously Presented) The compound of claim 49 wherein said fragment of (I) (b) differs from a peptide of (I) (a) by deletion of not more than 8 amino acids.

55. (Previously Presented) The compound of claim 49 wherein said fragment of (I) (b) differs from a peptide of (I) (a) by deletion of not more than 6 amino acids.

56. (Previously Presented) The compound of claim 49 wherein said fragment of (I) (b) differs from a peptide of (I) (a) by deletion of not more than 4 amino acids.

57. (Previously Presented) The compound of claim 49 wherein said fragment of (I) (b) differs from a peptide of (I) (a) by deletion of not more than 2 amino acids.

58. (Previously Presented) The compound of claim 49 that is a peptide of (I) (a) or (I) (b).

59. (Previously Presented) The compound of claim 49 that is a peptide of (I) (a).

60. (Previously Presented) The compound of claim 45 wherein said oligomer or polymer of (II) comprises at least two identical peptides according to (I).

61. (Previously Presented) The compound of claim 45 wherein said oligomer or polymer of (II) comprises at least four identical peptides according to (I).

62. (Previously Presented) The compound of claim 61 wherein the carrier is a lysine dendrimer backbone or bovine serum albumin.

63. (Previously Presented) The compound of claim 62 wherein the fragment of I(b) differs from a peptide of (I)(a) by deletion of not more than 2 amino acids, and the peptide of (I)(d) is at least 85% identical to a peptide of (I)(a).

64. (Previously Presented) The compound of claim 49 that is a peptide of (I)(a), (I)(b) or (I)(c).

65. (Previously Presented) The compound of claim 49 that is a peptide of (I)(a) or (I)(c).

66. (New) The compound of claim 49 that is a peptide of (I)(a), (I)(b), (I)(c) or (I)(d).

67. (New) The compound of claim 49 that is a peptide of (I)(a), (I)(b), (I)(c) or I(d1).

68. (New) The compound of claim 49 that is a peptide of (I)(a), (I)(b), (I)(c) or I(d2).

69. (New) The compound of claim 8, in soluble form.

70. (New) The compound of claim 8, which is not part of a combinatorial library.

71. (New) The compound of claim 8 wherein the peptide sequence of (I)(a) is SEQ ID NO:1, 7 or 40.

72. (New) The compound according to claim 71 wherein the carrier of (II) is a protein.

73. (New) The compound according to claim 71 wherein the carrier of (II) is bovine serum albumin.

74. (New) The compound according to claim 71 wherein the carrier of (II) is a lipophilic dendrimer, a micelle-like carrier formed by liphophilic derivatives, a starburst carbon chain polymer conjugate, or a ligand presenting assembly based on a derivative of diethylaminomethane.

75. (New) The compound according to claim 71 wherein (I) applies.

76. (New) The compound of claim 71 wherein the peptide of (I) (d) comprises a sequence at least 85% identical to a peptide of (I) (a).

77. (New) The compound of claim 71 wherein the peptide of (I) (d) comprises a sequence differing from a peptide of (I) (a) solely by a single substitution.

78. (New) The compound of claim 71 wherein the peptide of (I) (d) comprises a sequence which differs from a peptide of (I) (a) solely by one or more conservative substitutions, such being defined as

(1) replacement of an amino acid selected from the group consisting of Gly, Ala, Val, Leu and Ile with another amino acid of the same group,

(2) replacement of an amino acid selected from group consisting of Asp, Glu, Asn and Gln with another amino acid of the same group,

(3) replacement of an amino acid selected from the group consisting of Phe, Tyr, Trp, His, and Pro with another amino acid of the same group,

(4) replacement of an amino acid selected from the group consisting of Arg, Lys and His with another amino acid of the same group,

(5) replacement of Cys with an amino acid selected from the group consisting of Asp, Glu, Lys, Arg, His, Asn, Gln, Ser, Thr and Tyr,

(6) replacement of an amino acid selected from the group consisting of Pro, Ala, Gly, Ser and Tyr with another amino acid of the same group, and

(7) replacement of an amino acid selected from the group consisting of Leu, Ile, Val and Met with another amino acid of the same group.

79. (New) The composition of claim 71 wherein the

peptide of (I) (c) comprises a sequence differing from a peptide of (I) (a) solely by a single conservative substitution.

80. (New) The compound of claim 71 wherein said fragment of (I) (b) differs from a peptide of (I) (a) by deletion of not more than 8 amino acids.

81. (New) The compound of claim 71 wherein said fragment of (I) (b) differs from a peptide of (I) (a) by deletion of not more than 6 amino acids.

82. (New) The compound of claim 71 wherein said fragment of (I) (b) differs from a peptide of (I) (a) by deletion of not more than 4 amino acids.

83. (New) The compound of claim 71 wherein said fragment of (I) (b) differs from a peptide of (I) (a) by deletion of not more than 2 amino acids.

84. (New) The compound of claim 71 that is a peptide of (I) (a) or (I) (b).

85. (New) The compound of claim 71 that is a peptide of (I) (a).

86. (New) The compound of claim 71 wherein said oligomer or polymer of (II) comprises at least two identical peptides according to (I).

87. (New) The compound of claim 71 wherein said oligomer or polymer of (II) comprises at least four identical peptides according to (I).

88. (New) The compound of claim 71 wherein the carrier is a lysine dendrimer backbone or bovine serum albumin.

89. (New) The compound of claim 71 wherein the fragment of I(b) differs from a peptide of (I) (a) by deletion of not more than 2 amino acids, and the peptide of (I) (d) is at least 85% identical to a peptide of (I) (a).

90. (New) The compound of claim 71 that is a peptide of (I) (a), (I) (b) or (I) (c).

91. (New) The compound of claim 71 that is a peptide of

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(I) (a) or (I) (c).

92. (New) The compound of claim 71 that is a peptide of (I) (a), (I) (b), (I) (c) or (I) (d).

93. (New) The compound of claim 71 that is a peptide of (I) (a), (I) (b), (I) (c) or I(d1).

94. (New) The compound of claim 71 that is a peptide of (I) (a), (I) (b), (I) (c) or I(d2).